Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	4297	scanning adj probe adj microscop\$	US-PGPUB; USPAT; EPO; JPO; DERWENT; IBM_TDB	OR	ON.	2005/12/20 15:48
L2	39575	oligonucleotide near5 probe	US-PGPUB; USPAT; EPO; JPO; DERWENT; IBM_TDB	OR	ON	2005/12/20 15:48
L3	210	l1 and l2	US-PGPUB; USPAT; EPO; JPO; DERWENT; IBM_TDB	OR	ON	2005/12/20 15:48
L4	6	(scanning adj probe adj microscop\$) same (oligonucleotide adj probe)	US-PGPUB; USPAT; EPO; JPO; DERWENT; IBM_TDB	OR	ON	2005/12/20 15:49

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NEWS 2 "Ask CAS" for self-help around the clock NEWS 3 SEP 09 ACD predicted properties enhanced in REGISTRY/ZREGISTRY

NEWS 4 OCT 03 MATHDI removed from STN

NEWS 5 OCT 04 CA/CAplus-Canadian Intellectual Property Office (CIPO) added to core patent offices

NEWS 6 OCT 13 New CAS Information Use Policies Effective October 17, 2005

NEWS 7 OCT 17 STN(R) AnaVist(TM), Version 1.01, allows the export/download of CAplus documents for use in third-party analysis and visualization tools

NEWS 8 OCT 27 Free KWIC format extended in full-text databases

NEWS 9 OCT 27 DIOGENES content streamlined

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NEWS 15° DEC 14 2006 MeSH terms loaded for MEDLINE file segment of TOXCENTER

NEWS 16 DEC 14 CA/CAplus to be enhanced with updated IPC codes

NEWS 17 DEC 16 MARPATprev will be removed from STN on December 31, 2005

NEWS EXPRESS DECEMBER 02 CURRENT VERSION FOR WINDOWS IS V8.01, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0jc(JP), AND CURRENT DISCOVER FILE IS DATED 02 DECEMBER 2005. V8.0 USERS CAN OBTAIN THE UPGRADE TO V8.01 AT http://download.cas.org/express/v8.0-Discover/

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=> s (scanning(w)probe(w)microscop?)/bi,ab 176631 SCANNING/BI 151594 SCANNING/AB 217303 PROBE/BI 186564 PROBE/AB

525148 MICROSCOP?/BI 469129 MICROSCOP?/AB L1 3977 (SCANNING(W)PROBE(W)MICROSCOP?)/BI,AB

=> s (oligonucleotide#(5a)probe#)/bi,ab 79087 OLIGONUCLEOTIDE#/BI 60536

OLIGONUCLEOTIDE#/AB 305939 PROBE#/BI 264136 PROBE#/AB

L2 11882 (OLIGONUCLEOTIDE#(5A)PROBE#)/BI,AB

=> |1 and |2

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=> s l1 and l2

L3 4 L1 AND L2

=> d his

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L1 3977 S (SCANNING(W)PROBE(W)MICROSCOP?)/BI,AB

L2 11882 S (OLIGONUCLEOTIDE#(5A)PROBE#)/BI,AB

L3 4 \$ L1 AND L2

=> d |3 1-4 bib ab

L3 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:533800 CAPLUS DN 141:85095 TI Controlled alignment of nano-barcodes encoding specific information for ***scanning*** ***probe*** ***microscopy*** (SPM) reading IN Chan, Selena; Su, Xing; Yamakawa, Mineo PA USA SO U.S. Pat. Appl. Publ., 37 pp., Cont.-in-part of U.S. Ser. No. 251,152. CODEN: USXXCO DT Patent LA English KIND DATE FAN.CNT 3 PATENT NO. **APPLICATION** DATE -----PI US 2004126820 A1 20040701 US 2003-667004 20030919 US 2004058328 A1 20040325 US 2002-20020920 WO 2004038037 251152 A2 20040506 20030922 W: AE, AG, AL, AM, AT, WO 2003-US29726 AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,

MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2002-251152 A2 20020920 US 2003-667004
A 20030919

LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO,

ZM, ZW RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG,

OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL,

TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA,

KG, KZ, MD, RU, TJ, TM, AT, BE, BG,

FI, FR, GB, GR, HU, IE, IT, LU,

AB The methods, app. and compns. disclosed herein concern the detection, identification and/or sequencing of biomols., such as nucleic acids or proteins. In certain embodiments of the invention, coded probes comprising a probe mol. attached to one or more nano-barcodes may be allowed to bind to one or more target mols. After binding and sepn. from unbound coded probes, the bound coded probes may be aligned on a surface and analyzed by ***scanning*** ***probe*** ***microscopy*** . The nano-barcodes may be any mol. or complex that is distinguishable by SPM, such as carbon nanotubes, fullerenes, submicrometer metallic barcodes, nanoparticles or quantum dots. Where the ***probes*** ***oligonucleotides*** , adjacent coded ***probes*** hybridized to a target nucleic acid may be ligated together before alignment and SPM anal. Compns. comprising coded probes are also disclosed herein. Systems for biomol, anal, may comprise an SPM instrument and at least one coded probe attached to a

L3 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN AN 2004:371098 CAPLUS

DN 140:388197

surface.

SY, TJ, TM,

ZM, ZW, AM, AZ, BY,

CH, CY, CZ, DE, DK, EE, ES,

TI Controlled alignment of nano-barcodes encoding specific information for ***scanning*** ***probe***

microscopy (spm) reading

IN Chan, Selena; Su, Xing; Yamakawa, Mineo

PA Intel Corporation, USA

SO PCT Int. Appl., 63 pp. CODEN: PIXXD2

DT Patent

LA English

PI WO 2004038037 A2 20040506 WO 2003-US29726 20030922 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC,

EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG US 2004058328 A1 20040325 US 2002-251152 20020920 US 2004126820 A1 20040701 US 2003-667004 20030919

PRAI US 2002-251152 A 20020920 US 2003-667004 A 20030919

The methods, app. and compns. disclosed herein concern the detection, identification and/or sequencing of biomols., such as nucleic acids or proteins. In certain embodiments of the invention, coded probes comprising a probe mol. attached to one or more nano-barcodes may be allowed to bind to one or more target mols. After binding and sepn. from unbound coded probes, the bound coded probes may be aligned on a surface and analyzed by ***scanning*** ***probe*** ***microscopy*** . The nano-barcodes may be any mol. or complex that is distinguishable by SPM, such as carbon nanotubes, fullerenes, submicrometer metallic barcodes, nanoparticles or quantum dots. Where the ***probes*** are ***oligonucleotides*** , adjacent coded ***probes*** hybridized to a target nucleic acid may be ligated together before alignment and SPM anal. Compns. comprising coded probes are also disclosed herein. Systems for biomol. anal. may comprise an SPM instrument and at least one coded probe attached to a surface.

L3 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN AN 2004:252088 CAPLUS

DN 140:249737

TI Controlled alignment of nanobarcodes encoding specific information for ***scanning*** ***probe***

microscopy (SPM) reading

IN Chan, Selena; Su, Xing; Yamakawa, Mineo

PA USA

SO U.S. Pat. Appl. Publ., 17 pp. CODEN: USXXCO

DT Patent

LA English

PI US 2004058328 20040325 US 2002-251152 A1 20020920 WO 2004027095 A1 20040401 WO 2003-20030905 W: AE, AG, AL, AM, AT, AU, AZ, BA, US28082 BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, A1 20050622 EP 2003-752088 TG EP 1543152 20030905 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, NL, SE, MC, PT, CZ, EE, HU, SK US 2004126820 A1 20040701 US 2003-667004 20030919 WO 2004038037 A2 20040506 20030922 W: AE, AG, AL, AM, AT, WO 2003-US29726 AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,

CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG US 2005208554 20050922 US 2005-77577 20050311 PRAI US 2002-251152 WO 2003-US28082 Α 20020920 W 20030905 US 2003-667004 20030919 Α AB The methods, app. and compns. disclosed herein concern the detection, identification and/or sequencing of biomols., such as nucleic acids or proteins. In certain embodiments of the invention, coded probes comprising a probe mol. attached to one or more nanobarcodes may be allowed to bind to one or more target mols. After binding and sepn. from unbound coded probes, the bound coded probes may be aligned on a surface and analyzed by ***scanning*** ***probe*** ***microscopy*** . The nanobarcodes may be any mol. or complex that is distinguishable by SPM, such as carbon nanotubes, fullerenes, submicrometer metallic barcodes, nanoparticles or quantum dots. Where the ***probes*** are ***oligonucleotides*** , adjacent coded ***probes***
hybridized to a target nucleic acid may be ligated together before alignment and SPM anal. Compns. comprising coded probes are also disclosed herein. Systems for biomol, anal, may comprise an SPM instrument and at least one coded probe attached to a

L3 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1998:534997 CAPLUS

DN 129:226603

TI Detection of nucleic acids with ***scanning*** ***probe*** ***microscopy***

IN Hori, Kunio; Takahashi, Isao; Okada, Takao

PA Olympus Optical Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 13 pp. CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1 PATENT NO. KIND DATE APPLICATION DATE -----

PI JP 10215899 A2 19980818 JP 1997-25219 19970207 US 6194148 B1 20010227 US 1998-19931 19980206

PRAI JP 1997-25219 A 19970207

AB A simplified method for detecting target nucleic acids is described, which method comprises (1) heating the mixt. of a sample and ***oligonucleotide*** ***probes*** to allow denaturation; (2) lowering the temp. to allow hybridization between the target nucleic acids in the sample and the shape, structure, and length of the hybrids with the ***scanning*** ***probe*** ***microscopy*** that includes scanning tunneling microscopy and at. force microscopy.

=> d his

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L2 11882 S (OLIGONUCLEOTIDE#(5A)PROBE#)/BLAB

L3 4 S L1 AND L2 => log y

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